

Billing Code 4410-09-M

DEPARTMENT OF JUSTICE DRUG ENFORCEMENT ADMINISTRATION MANUFACTURER OF CONTROLLED SUBSTANCES NOTICE OF APPLICATION

Pursuant to § 1301.33(a), Title 21 of the Code of
Federal Regulations (CFR), this is notice that on
September 9, 2011, Johnson Matthey Inc., Custom
Pharmaceuticals Department, 2003 Nolte Drive, West
Deptford, New Jersey 08066-1742, made application by
renewal to the Drug Enforcement Administration (DEA) to be
registered as a bulk manufacturer of the following basic
classes of controlled substances:

Drug	Schedule
Gamma Hydroxybutyric Acid (2010)	I
Tetrahydrocannabinols (7370)	I
Dihydromorphine (9145)	I
Difenoxin (9168)	I
Propiram (9649)	I
Amphetamine (1100)	II
Methamphetamine (1105)	II

Drug	Schedule
Lisdexamfetamine (1205)	II
Methylphenidate (1724)	II
Nabilone (7379)	II
Cocaine (9041)	II
Codeine (9050)	II
Dihydrocodeine (9120)	II
Oxycodone (9143)	II
Hydromorphone (9150)	II
Diphenoxylate (9170)	II
Ecgonine (9180)	II
Hydrocodone (9193)	II
Meperidine (9230)	II
Methadone (9250)	II
Methadone intermediate (9254)	II
Morphine (9300)	II
Thebaine (9333)	II
Oxymorphone (9652)	II
Noroxymorphone (9668)	II
Alfentanil (9737)	II
Remifentanil (9739)	II
Sufentanil (9740)	II
Fentanyl (9801)	II

The company plans to manufacture the listed controlled substances in bulk for sale to its customers.

Any other such applicant, and any person who is presently registered with DEA to manufacture such substances, may file comments or objections to the issuance of the proposed registration pursuant to 21 CFR § 1301.33(a).

Any such written comments or objections should be addressed, in quintuplicate, to the Drug Enforcement Administration, Office of Diversion Control, Federal Register Representative (ODL), 8701 Morrissette Drive, Springfield, Virginia 22152; and must be filed no later than [insert date 60 days from date of publication].

Joseph T. Rannazzisi Deputy Assistant Administrator Office of Diversion Control Drug Enforcement Administration

DATED: November 18, 2011

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